Osteopontin Promotes Protective Antigenic Tolerance againstExperimental Allergic Airway Disease

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Abstract

In the context of inflammation, osteopontin (Opn) is known to promote effector responses, facilitating a proinflammatory envi-ronment; however, its role during antigenic tolerance induction is unknown. Using a mouse model of asthma, we investigated the role of Opn during antigenic tolerance induction and its effects on associated regulatory cellular populations prior to disease initiation. Our experiments demonstrate that Opn drives protective antigenic tolerance by inducing accumulation of IFN-b- producing plasmacytoid dendritic cells, as well as regulatory T cells, in mediastinal lymph nodes. We also show that, in the absence of TLR triggers, recombinant Opn, and particularly its SLAYGLR motif, directly induces IFN-b expression in Ag-primed plasmacytoid dendritic cells, which renders them extra protective against induction of allergic airway disease upon transfer into recipient mice. Lastly, we show that blockade of type I IFNR prevents antigenic tolerance induction against experimental allergic asthma. Overall, we unveil a new role for Opn in setting up a tolerogenic milieu boosting antigenic tolerance induction, thus leading to prevention of allergic airway inflammation. Our results provide insight for the future design of immunotherapies against allergic asthma.

Introduction

Mechanisms of central and peripheral tolerance are crucial for maintaining immune system homeostasis and preventing exaggerated immune responses to intrinsically harmless self- or foreign Ags. Failure of this mechanism could lead to the development of chronic inflammation, such as allergic asthma and autoimmune diseases. Because the incidence of allergic disease has risen dramatically, much effort has been put into determining the control mechanisms of peripheral tolerance to allergens in an attempt to find a treatment or prevention strategy for allergic disease. Allergic asthma is a disease caused by aberrant TH2 immune responses to inhaled allergens leading to eosinophilic airway inflammation, mucus hypersecretion, and variable airway obstruction (1). Regulatory T (Treg) cells are important suppressors of dysregulated TH2 responses to inhaled Ags, because constitutive or induced deficiency of these cells leads to severe asthmatic reactions (2, 3). Likewise, several groups have demonstrated that conventional dendritic cells (cDCs) and plasmacytoid dendritic cells (pDCs) are key regulators of TH2 responses in allergic airway inflammation (4–7). As in many processes of immunoregulation, cytokines like TGF-b1 and IL-10 are also important regulators of tolerance to inhaled Ags (3, 8, 9).
Osteopontin (Opn) is a cytokine expressed by immune cells, such as activated T cells and dendritic cells (DCs), as well as by non- immune cells, including tumor cells and stromal cells (10–12). In inflammatory conditions, Opn affects DC function (5, 13–18) and can drive TH1, TH2, and TH17 effector immune responses (5, 11, 13, 14, 19, 20). In contrast, Opn is constitutively expressed by a great variety of cells under noninflammatory conditions (12, 21– 23), but its physiological significance is largely unknown. For example, secreted Opn (sOpn) is expressed in the bone marrow (BM) and also upon inflammatory conditions, primarily in the form of thrombin-cleaved fragments (24, 25). Opn fragments have binding motifs for several integrins: the SLAYGLR motif spe- cifically interacts with integrins a4b1, a9b1, and a4b7, whereas the RGD motif interacts with the avb3, avb5, avb1, and a5b1 integrins (12, 26). In addition, the C-terminal half of Opn interacts with certain CD44 variants (12, 27). Thrombin cleavage of Opn reveals the otherwise cryptic SLAYGLR domain, and this modi- fication is vital for its interaction with a9b1 integrin (28).
Recent reports show that Opn is expressed in Foxp3^{+} Treg cells(29, 30), suggesting its possible role in immune regulation. In thisstudy, we test whether Opn affects tolerance induction during in- tranasal (i.n.) administration of endotoxin-free Ag. Our results un- veil a novel role for Opn as a tolerance enhancer against allergic airway disease, setting up an immunoregulatory milieu and poten-tiating CCR7-expressing pDC recruitment to the draining lymph nodes (dLNs). In addition, we reveal that, in the absence of pathogen-associated molecular patterns, sOpn, and specifically itsintegrin-binding SLAYGLR motif, induces low levels of IFN-b expression in Ag-primed pDCs. SLAYGLR-treated pDCs are highly efficient at suppressing allergic airway inflammation via type I IFN. Finally, we show that type I IFNs are crucial during antigenic tol- erance induction against allergic airway disease.

Materials and Methods

Mice

BALB/c, C57BL/6J (B6), OVA-specific TCR-transgenic (Tg) C.Cg-Tg (DO11.10)10Dlo/J (DO11.10), C.129P2(B6)-IL-10tm1Cgn/J, B6(Cg)-Il10tm1.1Karp (Il10GFP), B6.129-Ifnb1tm1Lky/J (IfnbEYFP), B6.129S2-*Ifnar1*^{tm1Agt}/Mmjax (*Ifnar1*$^{2/2}$), and B6-Tg (C-type lectin domain family 4, member C [CLEC4C]–heparin binding EGF like growth factor [HBEGF]) 956Cln/J (pDC-specific type II C-type lectin [BDCA2]-diphtheria toxin receptor–[DTR]) mice were purchased from The Jackson Laboratory (Bar Harbor, ME). B6-Tg Foxp3-DTR/EGFP(depletion of Treg cell [DEREG]) mice were provided by T.S. B6.129S6(Cg)-secreted phosphoprotein 1 (Spp1)^{tm1Blh}/J (*Spp1*$^{2/2}$) mice were kindly provided by Dr. L. Liaw (Maine Medical Center Research Institute, Scarborough ME). All mice used in this study were 8–10-wk-old females. Mice were housed at the Animal Facility of the Biomedical Research Foundation of the Academy of Athens and at the University Hospital Ghent (Ghent, Belgium). Use of mice in this studywas reviewed and approved by the Bioethics Committee of the Bio- medical Research Foundation of the Academy of Athens, the Veteri- narian Office of Attica, and the Animal Ethics Committee of Ghent University. All procedures were in accordance with the National In- stitutes of Health Statement of Compliance (Assurance) with Standards for Humane

Care and Use of Laboratory Animals (A5736–01) and with the European Union Directive 86/609/EEC for the protection of ani- mals used for experimental purposes.

In vivo experimental protocols

For tolerance induction, mice received 200 mg of EndoGrade OVA (Hyglos) i.n. in the presence of 2.5 mg recombinant Opn protein (rOpn) (R&D Systems), 72 ng of synthetic secreted $Opn_{134–153}$ fragments (frOpn; IVPTVDVPNGRGDSLAYGLR), or PBS (control). The RGD domain (Arg-Gly-Asp) of frOpn1 is scrambled to RAA (Arg-Ala-Ala). The SLAYGLR domain (Ser-Leu-Ala-Tyr-Gly-Leu-Arg) of frOpn2 is scrambled to LRAGLRS (Leu-Arg-Ala-Gly-Leu-Arg-Ser). frOpn3 has the RGD and SLAYGLR domains scrambled to RAA and LRAGLRS, respectively (Caslo Laboratory ApS). Opn oligopeptide fragments have been described previously (31). Myelin oligodendrocyte glycoprotein $(MOG)_{35–55}$ peptide (MEVGWYRSPFSRVVHLYRNGK; Caslo Laboratory ApS) was also used for tolerance induction (250 mg per mouse

i.n.). Mice were euthanized 36–40 h later and analyzed. In certain ex- periments, tolerance was induced prior to allergic airway disease in- duction: mice received 200 mg of EndoGrade OVA (i.n.) on days 22, 21, and 0 in the presence of 2.5 mg of rOpn or 72 ng of frOpn1-3 synthetic fragments (i.n.). Control mice received PBS. Allergic airway disease (asthma) was subsequently induced: on day 10, mice were immunized with chicken OVA Grade V (Sigma-Aldrich; 50 mg) in 0.2 ml of aluminum hy- droxide (alum) (Serva), followed by three challenges with 5% aerosolized OVA between days 16 and 18, as described (5, 15). Mice were euthanized 2 d after the last aerosol treatment (on day 20). DEREG mice received 1 mg of diphtheria toxin (DT; Sigma-Aldrich) (32) or PBS i.p. on days 23 and 22 (6 h prior to i.n. OVA administration). For the DT control group, non-Tg littermates were administered DT. For pDC depletion, mice received 225 mg of 120G8 pDC-depleting Ab (IgG2a; Dendritics, Lyon, France) (33) or an IgG2a isotype control Ab i.p. on days 26, 25, 24, and 23. As in Fig. 4A, mice received 200 mg of EndoGrade OVA i.n., along with 2.5 mg of rOpn, on days 22, 21 and 0 and were euthanized 7 d later. For pDC depletion in *BDCA2-DTR* mice, 120 ng of DT was administered i.p. on days 24 and 23 in the 7-d tolerance-induction protocol (as in Fig. 4A) (34). Efficient pDC depletion ($95%) from dLNs in both approaches was determined by FACS analysis (data not shown). For neutralization of IFNAR1, mice received
20 mg of a polyclonal affinity-purified neutralizing Ab to mouse
IFNAR1 (clone MAR1-5A3; eBioscience) or goat IgG isotype control (R&D Systems) i.p. 2 h before tolerance induction. For allergy in- duction, after IFNAR1 neutralization and tolerance, mice were im- munized with chicken OVA on day 15, and OVA challenges were performed between days 21 and 23.

Enhanced pause

Lung function was measured in mice 24 h after the final OVA challenge (day 19) by whole body plethysmography (Buxco Technologies) to calculate enhanced pause (Penh). Responses to inhaled methacholine (Sigma-Aldrich) at concentrations of 3–100 mg/ml were measured for 1 min, as previously described (5).

Analysis of bronchoalveolar lavage and lung histology

Bronchoalveolar lavage (BAL) harvesting and analysis were performed as previously described (35, 32). For histological analysis, paraffin-embedded (4-mm) lung sections were stained with H&E and quantified as previously described (35). Goblet cells were quantified in Periodic acid–Schiff (PAS)-stained lung sections (5). A semiquantitative scoring system was used to grade the area of lung infiltrated, as previously described (36). Goblet cells were counted on PAS-stained sections using an arbitrary scoring system, as previously described (36).

pDC generation from BM, cultures, and adoptive transfer

For pDC generation, BM cells were isolated and cultured with recombinant human Flt3 ligand, as described (37). On day 11, $7AAD^{2}$ $CD3^{2}CD19^{2}CD11c^{+}B220^{+}PDCA1^{+}Siglec\text{-}H^{+}$ pDCs were sorted to 98% purity using a FACSAria III, after enrichment with a CD11c MicroBeads Kit (Miltenyi

Biotec). Sorted pDCs from BM cultures were conditioned for 16 h with 100 mg/ml EndoGrade OVA or LoTox *Der- matophagoides pteronyssinus* allergen 1 (Derp1; Indoor Biotechnol- ogies) in the presence of 250 ng/ml rOpn, 18.2 ng/ml frOpn1 or frOpn3, or PBS. After the culture with OVA, pDCs were washed and analyzed or transferred i.v. via the tail vein (10^6 cells per mouse). Allergic asthma was induced in mice 7 d later. Mice were euthanized 2 d after the last OVA challenge (Figs. 7B, 10A).

Cell culture, proliferation, and cytokine analysis

Isolated dLN cells (2 3 10^5–10^6) were cultured with 125 mg/ml OVA (Sigma-Aldrich) for 48 h. We performed proliferation assays with thymi- dine incorporation, as previously described (38). For certain experiments, proliferation of cells was measured as the percentage of Edu^+ cells by FACS, using a Molecular Probes kit. For cytokine measurements, we used ELISA kits for IL-5 and IFN-g (both from BD Biosciences), IL-4 and IL-13 (both from R&D Systems), and IFN-b (BioLegend).

Flow cytometry

Freshly isolated live ($7AAD^2$; BD Biosciences) dLN cells and in vitro– derived BM cells were stained with combinations of fluorochrome- conjugated Abs to CD4–Pacific Blue or CD4-PE/Cy5 (clone GK 1.5); CD3-PE/Cy7, CD3-PE/Cy5, CD3-PE, or CD3-FITC (17A2); CD11c-PE/ Cy7, CD11c-FITC, or CD11c-Orange 605 (N418); CD11b-PE/Cy7 or CD11b-FITC (M1/70); B220-PE or B220-BV 510 (RA3-6B2); CCR7-PE
(4B12); Siglec-H–Pacific Blue or Siglec-H–FITC (551); PDCA-1–PE or PDCA-1–FITC (927); CD19-PE/Cy7 or CD19-PE/Cy5 (6D5); CD25-PE (PC61.5) (BioLegend); and T1ST2 (DIH9) (T1ST2-FITC [MD Biosciences] or T1ST2–PE [BioLegend]). For intranuclear staining of Foxp3, a permeabilization kit and Abs (Foxp3–Pacific Blue or Foxp3-PE/CyC5; clone FJK-16s) were used (eBioscience). Flow cytometric measurements were performed using an Attune Acoustic Focusing Cytometer (Applied Biosystems) and a FACSAria III (BD). FACS sorting of pDCs was per- formed using a FACSAria III. Data analysis was performed with FlowJo software (TreeStar).

Chemotaxis assay

Sorted pDCs from lymph nodes (LNs) and spleen of BALB/c mice were treated with rOpn (500 ng/ml) for 18–20 h and assayed for migration in response to chemokine CCL19 or CCL21 (200 ng/ml) (R&D Systems). The lower chambers of Transwell plates (QCM Chemotaxis Cell Migration Assay, 24- well [5 mm], colorimetric; Millipore) were filled with 500 ml of serum-free medium in the presence or absence of chemokines. DCs (10^5 cells per 200 ml) resuspended in serum-free medium were deposited in the upper chambers of the Transwell plates and allowed to migrate for 4 h at 37˚C in 5% CO_2. For each experiment, pooled total cells from spleen and inguinal and mesenteric LNs of mice (n = 8) were used, and pDCs were isolated by sorting.

Suppression assay and Treg cell induction in vitro

BM-derived pDCs, pretreated or not with OVA and/or rOpn, were cultured with naive DO11.10 $CD4^+$ T cells for 3 d at a 1:5 ratio. T cells were harvested and cultured in the presence of 1 ng/ml recombinant mouse IL-2 (R&D Systems) for an additional 7 d. Suppressive activity was assayed on 10^5 freshly purified CFSE (Invitrogen)-labeled DO11.10 $CD4^+$ T cells stimulated with 10^4 irradiated BALB/c splenocytes, with 1 mg/ml OVA 323–339 peptide (Caslo Laboratory ApS), in the presence or absence of 10^5 DC-stimulated T cells. CFSE uptake was assayed 7 d later (Fig. 7A).

Quantitative real-time PCR analysis

Total RNA was extracted from cells isolated from dLNs with anti– mPDCA-1 and CD11c MicroBeads (Miltenyi Biotec) and further purified with FACS sorting. cDNA synthesis was performed as described (31). Primers were designed using the Primer3 program and are shown in Supplemental Table I (MWG Eurofins). *Hprt*, *Foxp3*, *Il-10*, *Il27p28*, and *Spp1* primers were described previously (31). Real-time PCR was performed and analyzed as previously described (31). The reference gene used for real-time PCR analysis was *Hprt*.

Statistical analysis

Data were analyzed using Prism 7 software (GraphPad). The two-tailed Student *t* test was used for statistical analyses of two-group comparisons. Multigroup comparisons were performed using two-way ANOVA, followed by the Bonferroni correction for the multiplicity of tests. All results are presented as mean 6 SEM. In all experiments, statistical significance was defined as $*p \# 0.0332$, $**p \# 0.0021$, $***p \# 0.0002$, and $****p$, 0.0001.

Results

Opn boosts antigenic tolerance leading to increased protection from allergic airway disease

To address whether Opn plays a role in tolerance induction, mice received endotoxin-free chicken OVA, together with endotoxin-free rOpn or PBS control i.n., and protection against disease was assessed using a well-established model of allergic asthma (39) (Fig. 1A). OVA-tolerized mice showed a significant decrease in BAL total cells and eosinophils, as well as lung tissue inflammatory scores and mucus secretion (Fig. 1B, 1C), compared with nontolerized mice. In addition, Penh in OVA-tolerized mice was significantly reduced (Fig. 1D). Importantly, in OVA/rOpn- tolerized mice, numbers of eosinophils in BAL (Fig. 1B), airway hyperresponsiveness (AHR) (Fig. 1D), lung leukocytic infiltration, and mucus secretion (Fig. 1C) were further reduced compared with OVA-tolerized mice. Further- more, OVA/rOpn tolerization resulted in significantly reduced levels of IL-4, IL-13, and IFN-g in BAL, as well as a reduction in OVA-specific responses in mediastinal lymph node (mLN) cell cultures and in T_H cell proliferation, compared with OVA tolerization (Fig. 1E–G). Studies from our group (5, 15) and other investigators (40–42) have shown that there are high levels of IFN-g production in allergic airway disease. The above findings indicated that administration of Opn, along with OVA, promotes enhanced toler-ance, conferring significant protection from disease development.

We next investigated whether Opn deficiency had any effect on tolerance induction. Tolerance induction in $Spp1^{2/2}$ mice was not as effective as in *Spp1* mice, because we did not note a significant change in the number of eosinophils or total cell numbers in BAL (Fig. 2A). In accordance, tolerogenic i.n. OVA administration in $Spp1^{2/2}$ mice could not efficiently dampen the inflammation and mucus secretion in the lung, whereas it was very efficient in $Spp1^{+/+}$ mice (Fig. 2B). T_H2 cytokine production by OVA–stimulated mLN cells was also lower in OVA-tolerized $Spp1^{2/2}$ mice compared with PBS-treated $Spp1^{2/2}$ mice (Fig. 2C). OVA-stimulated mLN cells from OVA-tolerized $Spp1^{+/+}$ mice produced ~50% lower levels of T_H2 cytokines compared with cells from PBS-treated $Spp1^{+/+}$ mice, whereas OVA-tolerized $Spp1^{2/2}$ mice had a smaller reduction in cytokine levels compared with $Spp1^{+/+}$ mice (Fig. 2C). The reduction in IFN-g levels was similar among groups (Fig. 2C). Finally, OVA tolerization of $Spp1^{2/2}$ mice resulted in a smaller decrease in the percentages of $CD3^+$ T proliferating cells in cultures of mLN cells compared with those from $Spp1^{+/+}$ OVA-treated mice (Fig. 2D). The above results strongly support that tolerance induc-tion is more effective in the presence of Opn.

Opn administration along with Ag increases accumulation of tolerogenic pDCs

Migratory nonlymphoid tissue DCs transporting Ags to LNs are involved in promoting tolerance to self-antigens in the steady-state. Because pDCs constitute a tolerogenic DC subset (43–47), and Opn has a dual role in the recruitment of DC subsets (5), we ana- lyzed pDC numbers in dLNs of OVA-tolerized mice and PBS-treated controls (Fig. 3A). Numbers of dLN pDCs were increased in the OVA-treated group compared with the PBS control group (Fig. 3B). Mice treated with OVA/rOpn had increased percentages and sig- nificantly higher total numbers of pDCs in the dLNs at 36–40 h following treatment compared with OVA treatment (Fig. 3B). In contrast, cDC numbers in the dLNs of OVA/rOpn mice were significantly reduced and percentages were lower compared with OVA treatment alone (Fig. 3B). Therefore, administration of Opn during tolerance induction affects the balance of DC subsets in dLNs by increasing the numbers of pDCs and re- ducing the numbers of cDCs.

The importance of Opn for DC subset recruitment in tolerance was also demonstrated using Opn-deficient mice. Antigenic tol- erance induction in $Spp1^{2/2}$ mice resulted in a significant re- duction in pDC numbers in their dLNs compared with $Spp1^{+/+}$ mice (Supplemental Fig. 1). At the same time, cDC numbers were significantly increased in dLNs of $Spp1^{2/2}$ mice (Supplemental Fig. 1). Conclusively,

these results show that tolerogenic admin- istration of OVA in an Opn-efficient microenvironment enhances pDC accumulation in dLNs.

Administration of Opn during tolerance induction regulates CCR7 expression affecting DC subset homing to dLNs

To explore the reason for the increased numbers of pDCs in dLNs of Opn-treated mice, we investigated the effect of Opn on $CCR7^+$ pDCs. CCR7 is a chemokine receptor responsible for homing of DCs to dLNs (48–50). We quantified $CCR7^+$ DC subsets in pe- ripheral blood 36 h following OVA/rOpn tolerogenic administra- tion (Fig. 3A). The percentages of $CCR7^+$ pDCs per total pDC numbers were significantly elevated in the peripheral blood of mice treated with rOpn during tolerance induction compared with PBS-treated mice (Fig. 3C). The percentages of $CCR7^+$ cDCs per total cDC numbers originating from the peripheral blood of mice that had been administered OVA/rOpn were significantly de- creased compared with OVA-administered mice (Fig. 3C). Our results demonstrate that Opn leads to enhanced migratory $CCR7^+$ pDC percentages per total pDCs in the blood.

We also found that OVA/rOpn-administered mice had signifi- cantly higher levels of *Ccl19* and *Ccl21* (encoding CCL19 and CCL21 chemokines that bind to CCR7) expression in their dLNs compared with OVA-administered mice (Fig. 3D), possibly attracting the increased numbers of $CCR7^+$ pDCs to dLNs. Indeed, in vitro transmigration assays showed that sorted pDCs from naive mice pretreated with rOpn had a 2-fold increase in CCL19-induced chemotaxis compared with PBS-treated pDCs (Fig. 3E). The above findings demonstrate that, during toler- ance induction, Opn can differentially regulate the percentages of $CCR7^+$ DC subsets, thereby affecting their chemotactic migration to dLNs in favor of pDCs. Moreover, the observed increased gene expression of the CCR7 ligands CCL19 and CCL21 provide an extra explanation for the rOpn-mediated migration of pDCs to dLNs.

Opn enhances pDC-dependent $Foxp3^+$ Treg cell accumulation and promotes an immunoregulatory milieu in dLNs

Because Ag administration for tolerance induction leads to gen- eration of Treg cells (51), we investigated whether Opn had an effect early on this process. We administered OVA with rOpn (OVA/rOpn) or PBS i.n. to mice for three consecutive days, and mice were analyzed 1 wk later (Fig. 4A). OVA/rOpn-treated mice showed a nearly 2-fold increase in the percentages, as well as in the total numbers, of $CD4^+Foxp3^+$ Treg cells in dLNs compared with OVA-treated mice (Fig. 4B). Increased numbers of $Foxp3^+$ Treg cells in OVA/rOpn-tolerized mice were accompanied by significantly enhanced *Foxp3* expression in dLN cells (Fig. 4C). In addition, OVA/rOpn treatment induced a significant increase in the expression levels of immunoregulatory molecules, such as *Il10*, *Ido*, *Tgfb1*, *Fgl2*, and *Il27*, in dLNs compared with OVA treatment (Fig. 4C). Conclusively, Opn administration, along with Ag, promotes $Foxp3^+$ Treg cell accumulation, resulting in a highly tolerogenic microenvironment in the dLN.

To directly test whether increased numbers of $Foxp3^+$ Treg cells mediate the observed rOpn promotion of tolerance, we depleted $Foxp3^+$ Treg cells prior to tolerance induction using DEREG mice (52) (Fig. 5A). Induction of tolerance with rOpn could not protect Treg-depleted mice, which exhibited increased allergic responses, as demonstrated by increased eosinophilia and lymphocytosis in the BAL, increased T cell proliferation, elevated lung inflammation with enhanced mucus production, and increased OVA-specific cytokine responses compared with Treg-sufficient mice (Fig. 5B–E).

To address whether the Opn-mediated increase in pDC numbers was responsible for the observed accumulation of $Foxp3^+$ Treg cells, we depleted pDCs by administering 120G8 pDC-depleting Ab (33) prior to induction of tolerance and examined the accu- mulation of $Foxp3^+$ Treg cells in dLNs. This depletion led to a significant reduction in $CD4^+Foxp3^+$ Treg cell numbers compared with OVA/rOpn administration in non–pDC-depleted mice (Fig. 6A), resulting in exacerbated allergic airway disease (Fig. 6B). We observed similar effects of rOpn on Treg cellularity in LNs after in vivo depletion of pDCs in *BDCA2-DTR* Tg mice

(34) (Supplemental Fig. 2). Thus, Opn-mediated pDC accumula- tion is responsible, at least in part, for the higher numbers of Treg cells in dLNs. In addition, dLN cells from OVA/rOpn-treated

pDC-depleted mice were significantly less capable of suppress- ing OVA-specific T cell proliferation (Fig. 6C) compared with dLN cells from OVA/rOpn-treated non–pDC-depleted mice, reaching an even lower suppressive ability than that of OVA- treated pDC-depleted dLN cells. Thus, OVA/rOpn treatment increased the numbers of pDCs, favoring the accumulation of $Foxp3^+$ Treg cells that are crucial for tolerance maintenance.

Opn treatment of pDCs increases their suppressive activity against allergic airway disease

We tested the suppressive function of in vitro OVA/rOpn-treated BM-derived pDCs by coculturing them with $CD4^+$ T cells. T cells obtained from OVA/rOpn-treated pDC cultures signifi- cantly suppressed the responses of DO11.10 T cells to OVA compared with T cells obtained from OVA-treated pDC cultures (Fig. 7A). We also adoptively transferred BM-derived pDCs, which were preconditioned in vitro with OVA or OVA/rOpn, into recipient mice prior to induction of allergic airway inflammation (Fig. 7B).

Control mice received PBS-treated pDCs. Total and eosinophil cell numbers, as well as lung tissue inflammatory scores and mucus secretion, were lower in BAL of OVA/rOpn-treated pDC recipient mice compared with OVA-treated pDC recipient mice (Fig. 7B, 7C). Furthermore, OVA/rOpn-treated pDC recipients had significantly reduced OVA-specific responses compared with OVA-treated pDC recipients (Fig. 7D). Thus, OVA/rOpn treatment of pDCs rendered them more regulatory, indicating that Opn af- fects the intrinsic tolerogenic function of pDCs.

Opn SLAYGLR motif is responsible for pDC recruitment and effective protection from allergic disease

RNA expression analysis revealed that, during tolerance in- duction, mLN pDCs have increased expression of integrins avb3, a4b1, and a9b1 (data not shown). Because the SLAYGLR motif of Opn interacts with a4b1, a4b7, and a9b1 integrins (12), and the RGD motif interacts with the avb3 integrin (11, 12), we asked which Opn domain is responsible for the observed effects on pDCs during tolerance induction. Thus, synthetic $frOpn_{134–153}$, containing the RGD and the SLAYGLR integrin binding motifs, either intact or scrambled(31), were used with OVA to induce tolerance in mice, allowing us to unveil the involvement of the different integrin-binding motifs of Opn in pDC recruitment (Fig. 8A). $frOpn_{134–153}$ rep- resents the thrombin cleaved fragment of Opn that reveals the otherwise cryptic domain SLAYGLR (28). OVA/frOpn1, con-taining an intact SLAYGLR motif but a scrambled RGD motif, induced the accumulation of higher numbers of pDCs in LNs compared with OVA and OVA/rOpn (Fig. 8A). Conversely, frOpn2, which contains an intact RGD, but a scrambled SLAYGLR motif, induced a reduction in pDC numbers in dLNs compared with all other treatments (Fig. 8A). When both motifs were scrambled, as in the case of frOpn3, the numbers and percentages of pDCs were not significantly altered compared with OVA (Fig. 8A, Supplemental Fig. 3A). OVA/frOpn1 in- duced higher percentages of pDCs in dLNs compared with OVA and OVA/frOpn3 (Supplemental Fig. 3A). These data revealed that the Opn SLAYGLR motif is responsible for pDC accumulation in dLNs. These data also reveal that the RGD motif suppresses this accumulation.

To examine the in vivo tolerogenic potential of frOpn1 in al- lergic disease, mice were treated with OVA, together with frOpn1 or frOpn3 for control, before the induction of allergic airway inflammation (Fig. 8B). OVA/frOpn1 was more successful than OVA/frOpn3 in promoting tolerance and, thus, protecting mice against allergic airway inflammation, as demonstrated by lower Penh values (Fig. 8C). BAL eosinophilia, as well as inflamma- tory score and mucus staining, was also dampened after OVA/ frOpn1 treatment (Fig. 8D). In addition, treatment with OVA/ frOpn1 led to reduced levels of OVA-specific responses (Fig. 8E) and numbers of $T1ST2^+CD4^+$ T cells in dLNs (Fig. 8D). Collectively, this dataset indicates that the Opn SLAYGLR motif significantly boosts tolerance.

*Opn SLAYGLR motif induces IFN-*b *expression in pDCs*

We further asked whether OVA/frOpn1 treatment affects in vivo pDC immune profile 36–40 h after tolerance induction (Fig. 3A). In vitro, OVA/frOpn1 conditioning of pDCs promoted a 2-fold increase in *Ifnb* mRNA expression at 16 h (Fig. 9A). The observed difference in *Ifnb* expression was also reflected in the amount of IFN-b secreted, which was 5-fold higher in OVA/ frOpn1-conditioned pDCs compared with OVA/frOpn3- conditioned pDCs (Fig. 9B). House dust mite endotoxin-free Derp1 was also used to test whether the increased IFN-b response could be observed using another allergen. Indeed, Derp1/frOpn1 in vitro–conditioning of pDCs resulted in higher *Ifnb* expression

compared with Derp1/frOpn3 (Fig. 9C). We also measured higher levels of *Ifnb* expression in OVA/rOpn- conditioned pDCs compared with OVA/PBS-conditioned pDCs (Supplemental Fig. 3B). RNA expression analysis using mLN pDCs isolated from mice treated with OVA versus OVA/rOpn showed differential expression of *Ifnb* compared to mice treated with OVA (data not shown). We found that, indeed, *Ifnb* expression was upregulated in mLN pDCs isolated from mice treated with OVA/frOpn1 compared with those isolated from mice treated with OVA/frOpn3 (Fig. 9E). To confirm the changes in IFN-b expression in vivo, we used IFN-b^{EYFP} reporter mice. OVA/frOpn1-tolerized mice had significantly increased numbers of IFN-b^{EYFP+} pDCs and higher IFN-b expression per pDC (mean fluorescence intensity [MFI]) compared with OVA/frOpn3-tolerized mice (Fig. 9D). To test whether this effect is relevant to the Ag used, we tolerized IFN-b^{EYFP} mice with a self-antigen, $MOG_{35–55}$, with frOpn1 or frOpn3. $MOG_{35–55}$/frOpn1 administration to IFN-b^{EYFP} mice resulted in significantly increased accumulation of IFN-b^{EYFP+} pDCs and IFN-b expression per pDC (MFI) in LNs (Fig. 9F). However, the increase was modest compared with Opn/OVA, possibly because MOG is a peptide. These results suggest that, under tolerogenic conditions, the Opn SLAYGLR motif can boost *Ifnb* expression in pDCs through a mechanism in- dependent of the Ag used.

*Opn/SLAYGLR-induced IFN-*b *in pDCs is protective against allergic disease*

pDCs were primed in vitro with endotoxin-free OVA and frOpn1 or frOpn3 and transferred into mice prior to induction of allergic airway inflammation (Fig. 10A). Recipient mice were *Ifnar1*$^{+/+}$ or *Ifnar1*$^{2/2}$. pDCs treated with frOpn1 were significantly more potent at protecting *Ifnar1*$^{+/+}$ mice against allergic airway in- flammation, as demonstrated by reduced lung airway inflamma- tion, BAL eosinophilia, and goblet cell hyperplasia, compared with the control and OVA groups (Fig. 10A). Similar enhanced protection was also observed when pDCs were treated with rOpn and adoptively transferred into *Ifnar1*$^{+/+}$ mice (Fig. 7B–D). Ac- tually, type I IFN produced by pDCs was crucial in maintaining this protective effect, because *Ifnar1*$^{2/2}$ recipient mice were not protected against allergic asthma (Fig. 10A). These results dem- onstrate that the integrin-binding SLAYGLR domain of Opn renders pDCs more regulatory through type I IFN production in the context of allergic airway inflammation. In fact, pDCs cannot preserve their regulatory function in recipients that lack expres- sion of type I IFNR. Finally, neutralization of IFNAR1 during the OVA tolerance-induction phase (Fig. 10B) resulted in reduced protection from allergic airway disease concomitant with in- creased eosinophilia in BAL (Fig. 10C) and increased OVA- specific responses (Fig. 10D). This indicates that type I IFN is absolutely necessary for effective antigenic tolerance induction in this context.

Discussion

In this article, we unveil a new role for Opn, and particularly its SLAYGLR motif, in setting up a tolerogenic milieu driving anti- genic tolerance induction and, thus, leading to protection from allergic airway inflammation. The respiratory mucosa is constantly being exposed to a myriad of nonpathogenic environmental Ags. To protect against the im- munopathological consequences of this constant stimulus, a default low noninflammatory TH2 mechanism and/or a T cell–mediated tolerance mechanism is activated (53). The mechanism underlying these processes is not fully understood. In this study, we initially demonstrate that Opn induces tolerance by tilting the pDC/cDC balance in favor of anti-inflammatory pDCs, via differentially regulating their CCR7 expression. CCR7-dependent homing of DCs into dLNs is required for the induction of tolerance (50).

Opn promotes T_H2 effector responses when administered during the allergen-sensitization phase (5). In contrast, we have previ- ously shown that Opn inhibits T_H2 responses during allergen challenge, and administration of rOpn during that phase is sup- pressive for experimental allergic asthma (5). In this study, we explore the role of Opn during a different phase, which is when Ag is administered during tolerance induction, prior to sensitiza- tion. Thus, Opn is administered i.n., prior to the sensitization phase, along with the allergen (antigen), to test its effects upon antigenic tolerance induction. Together, our results from this study, as well as from a previous study (5), show that adminis- tration of Opn during antigenic tolerance induction and during antigenic challenge is protective and, thus, can be used as a pre- ventive or a therapeutic agent against allergic airway disease.

Regulatory cytokines, such as *Tgfb1*, *Il10*, and *Fgl2*, and mol- ecules, such as *Ido* (54), were substantially upregulated in our Opn-driven tolerance model, creating an immunoregulatory microenvironment in the dLN. This tolerogenic milieu was accompanied by the accumulation of $Foxp3^+$ Treg cells, which are very important for induction of tolerance (51). Mice that had en- hanced Treg cell numbers due to rOpn administration showed suppressed effector responses and were protected from allergic disease. When $Foxp3^+$ Treg cells were depleted, Opn-mediated tolerance boosting was abolished. In light of its well-established proinflammatory function (10–12), it was unexpected that Opn could be an inducer of Treg cell accumulation in vivo. In the absence of pDCs, rOpn administration was no longer capable of inducing increased numbers of $Foxp3^+$ Treg cells in dLNs, demonstrating that rOpn affects Treg cell numbers primarily through its impact on pDCs. In contrast, after tolerance induction without the addition of Opn, Treg cell numbers do not appear to depend on pDCs, because pDC depletion did not reduce their numbers. These results show that rOpn conditions pDCs to enhance Treg cell accumulation. Functional flexibility and fostering of Treg cell responses are typical features of pDC involve-ment in tolerance (4, 46), as also revealed by our studies.

Thrombin cleavage of Opn occurs during homeostatic and inflam- matory conditions (24, 25), and conformational changes after thrombin cleavage result in higher-affinity binding to certain receptors (26). In fact, the cryptic SLAYGLR motif is revealed upon thrombin cleav-age of Opn, which is vital for its interaction with a9b1 integrin (28). Because the SLAYGLR motif of Opn protein interacts with a4b1, a4b7, and a9b1 integrins (12), and the RGD motif in- teracts with the avb3 integrin (11, 12), we used synthetic $frOpn_{134–153}$, containing both integrin binding motifs (intact or scrambled) (31). Thrombin cleavage of Opn also produces a fragment containing the C-terminal half of Opn, which interacts with certain CD44 variants (12, 27). In our settings, as well as in other settings (31, 55, 56), $frOpn_{134–153}$ has a greater efficiency compared with full-length Opn. Thus, it is possible that inter- action of Opn with CD44 interferes with certain Opn effects, such as pDC recruitment. Future investigations will elucidate the role of Opn–CD44 interactions in tolerance induction.

Under endotoxin-free conditions, we demonstrated that the SLAYGLR domain of Opn (frOpn1) enhances IFN-b expression in pDCs exposed to protein or peptide Ag. Accordingly, we showed that adoptively transferred OVA/frOpn1-treated pDCs were more efficient at suppressing allergic airway inflammation in recipient mice. This regulatory function was indeed mediated by the ob- served upregulated production of IFN-b by pDCs, because *Ifnar1*$^{2/2}$ recipient mice were not protected against allergic asthma. During the course of allergic airway inflammation, pDCs, as well as their type I IFN production, are suppressive for disease (4, 57–59). Our studies reveal that pDCs expressing higher IFN-b levels due to exposure to Opn are important for protective toler- ance prior to disease induction. Importantly, our experiments point to a crucial role for type I IFNs during the induction of efficient

antiallergic antigenic tolerance. Accordingly, IFNAR signaling promotes Treg cell development and function understress conditions (60).

Opn administered during tolerance also resulted in a remarkableupregulation of the ligands of CCR7, CCL19, and CCL21 in dLNs,reminiscent of that observed in tumors (61). Specifically, high CCL21 expression by melanoma in mice was associated with an immunotolerant microenvironment, which included the induction oflymphoid-like reticular stromal networks, an altered cytokine mi- lieu, and, most importantly, the recruitment of regulatory leukocyte populations (61). High Opn expression is prevalent in many types ofmalignancy (62); although Opn is considered proinflammatory, most of these tumors escape immune surveillance. Thus, it is pos- sible that Opn and its mediated induction of CCL21 could also be involved in suppression of antitumor immunity.

In this article, we demonstrate that the SLAYGLR motif of sOpnenhances regulatory mechanisms when administered together withendotoxin-free Ag in a tolerogenic context. In this respect, Opn hasa unique effect on immunity, differing substantially from its effects in the presence of danger signals. The SLAYGLR motif of Opn interacts with integrins (12). Addressing which integrin mediates the tolerogenic effects of the SLAYGLR motif of Opn on Ag- loaded pDCs, and primarily the induction of IFN-b, will assistin the design of therapies targeting tolerance in allergy. Finally, our results point to novel effects of Opn on Foxp3$^+$ Treg cells thatremain to be explored.

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Disclosures
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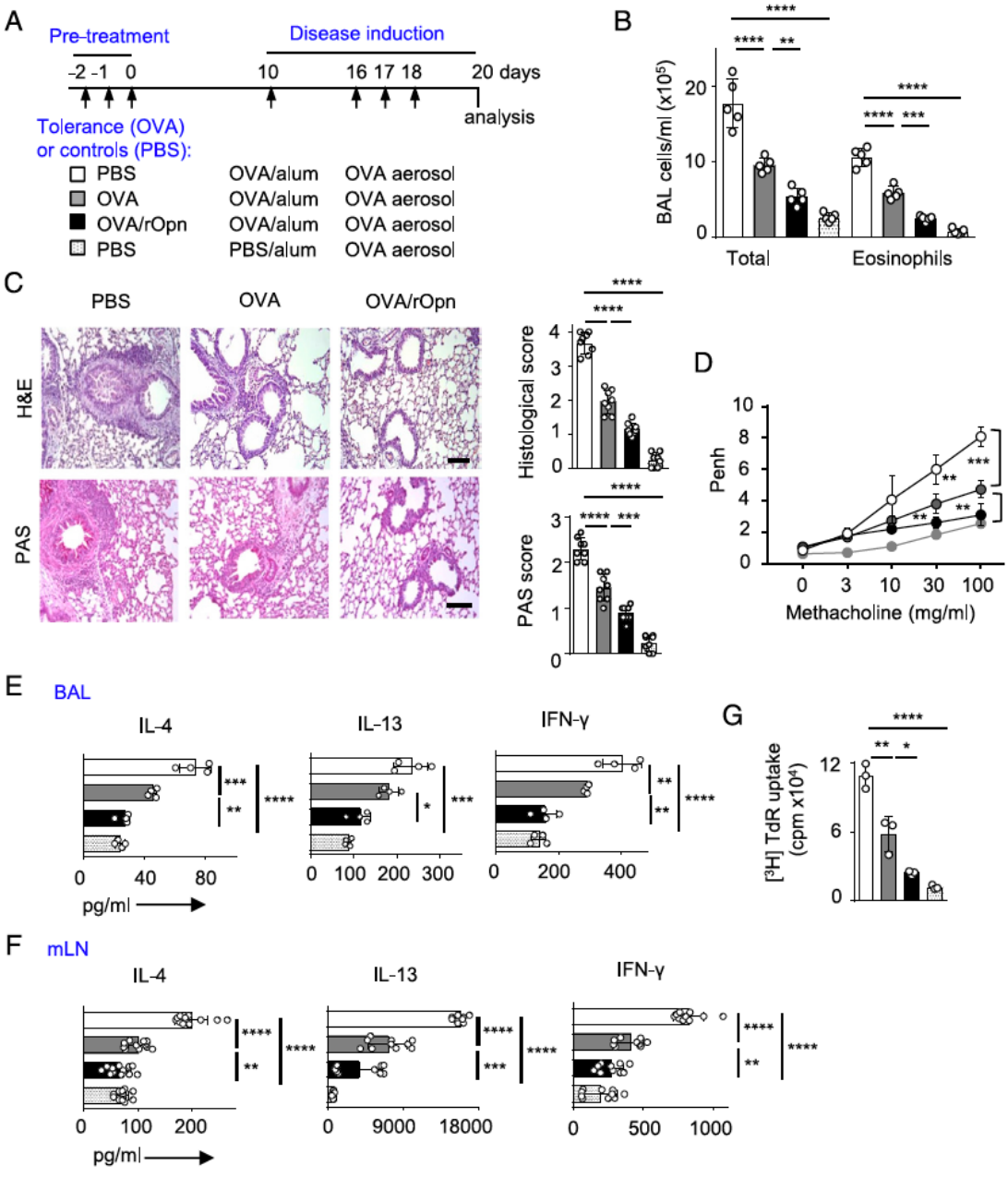


FIGURE 1. Opn administration boosts antigenic tolerance, leading to increased protection from allergic airway disease. (A) Experimental protocol used forendotoxin free-tolerance (OVA administration) induction, followed by allergic asthma induction. Controls were mice pretreated with PBS (no tolerance) and sub-sequent allergic airway disease induction (white bars). Controls for allergic airway disease were mice pretreated and sensitized with PBS. (B) Total cell counts (inBAL) and eosinophils. (C) Representative photomicrographs of H&E- and PAS-stained lung sections and histological scores. Scale bars, 100 mm. (D) AHR (depicted as Penh) on day 19. Levels of IL-4, IL-13, and IFN-g in BAL (E) and in the supernatant of mLN cells (F) stimulated ex vivo with OVA. (G) [^{3}H]Thymidineincorporation of mLN cells stimulated ex vivo with OVA. The same numbers of pooled mLN cells were cultured in 3, 4, 10, or 12 different wells per group in (E)–(G). Data are mean 6 SEM (n = 8 mice per group), one representative of three independent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p ¸ 0.0001.

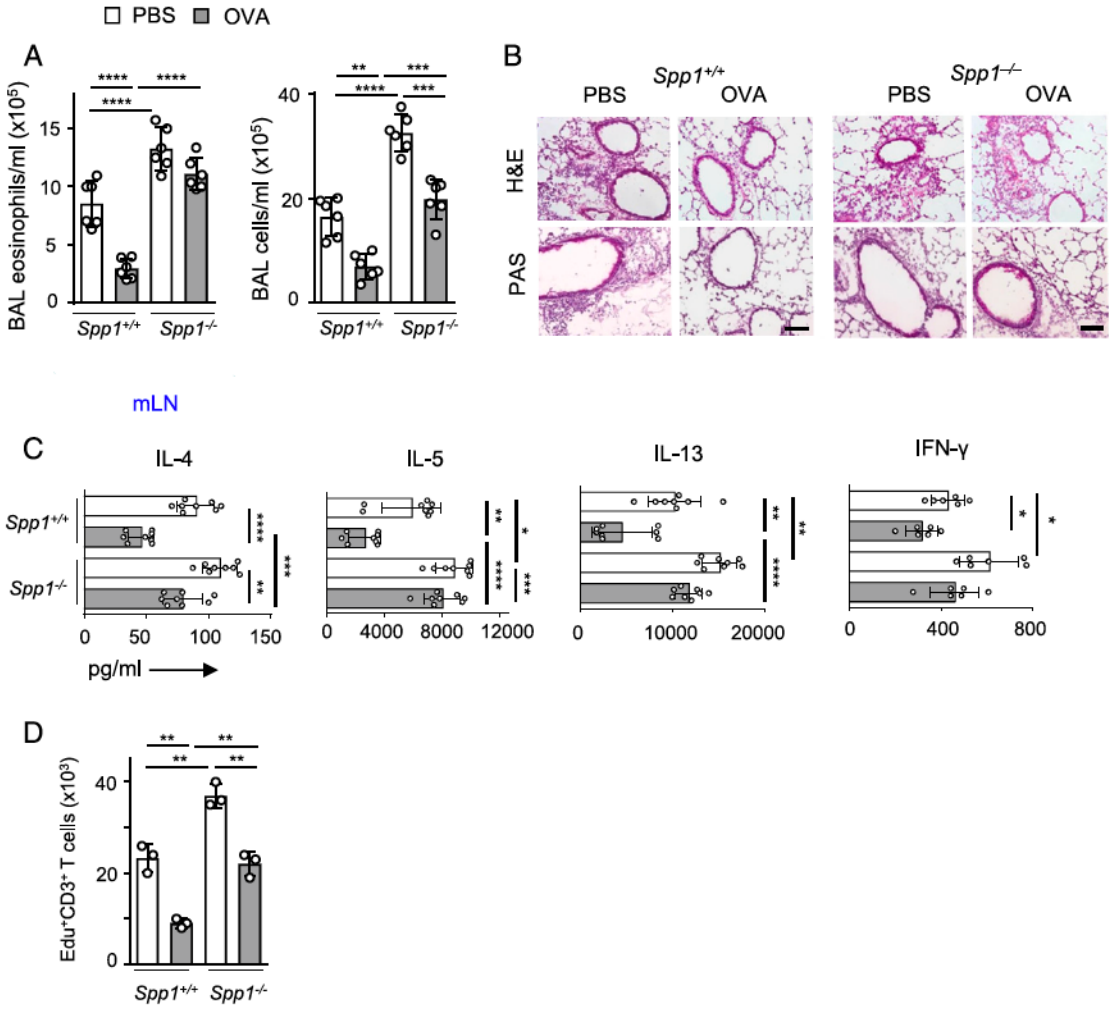


FIGURE 2. *Spp1*$^{2/2}$ mice are more resistant to tolerance induction. *Spp1*$^{2/2}$ or *Spp1*$^{+/+}$ mice received three doses of endotoxin-free OVA or PBS i.n. on days22, 21, and 0 (as in Fig. 1A). On day 10, mice were sensitized i.p. with OVA in alum and were subsequently challenged through the airways with aerosolized OVA between days 16 and 18. Analysis was performed on day 20. (A) Differential eosinophil and total cell counts in BAL. (B) Lung inflammation in H&E-stained sections and mucus secretion in PAS-stained sections from PBS- and OVA-treated *Spp1*$^{+/+}$ and *Spp1*$^{2/2}$ mice. Scale bar, 100 mm. Levels of IL-4, IL-5, IL-13, and IFN-g in supernatants of OVA-stimulated dLN cells (C) and percentages of Edu^{+}CD3^{+}CD4^{+} T cells in OVA-stimulated dLN cells (D) in both micegroups. The same numbers of pooled mLN cells were cultured in three, six, or eight different wells per group in (C) and (D). Data are mean 6 SEM (n = 6 mice per group), one representative of three independent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.

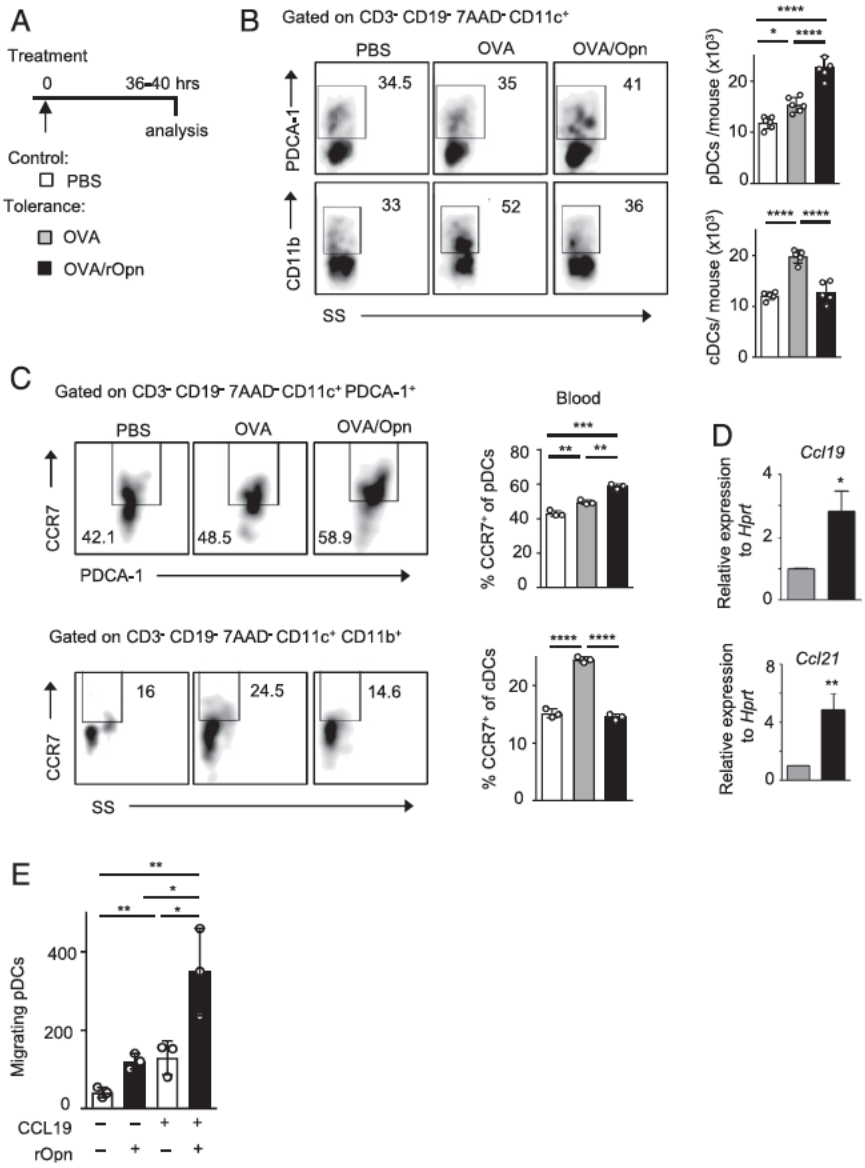


FIGURE 3. Opn-induced accumulation of pDCs in dLN and differential CCR7 expres- sion. Treatment of mice with endotoxin-free OVA (tolerance induction), together with endotoxin-free rOpn (A) or treatment of mice with PBS without OVA for control (B–D). Controls (without tolerance) were PBS- treated mice (white bars). (B–D) 7AAD2 CD32CD192CD11c+PDCA-1+CCR7+ pDCs and 7AAD2CD32CD192CD11c+CD11b+ CCR7$^+$ cDCs quantified in dLNs by flow cytometry. (B) Representative percentages in flow cytometric plots and numbers of dLN pDCs (upper panels) and cDCs (lower pan-els). (C) Representative flow cytometric plots and percentages of peripheral blood CCR7$^+$ pDCs and cDCs among total pDCs and cDCs, respectively. (D) Relative expression to *Hprt* of *Ccl19* and *Ccl21* in dLNs of tolerized mice. Data are mean 6 SEM (n = 10 mice per group; cDNAs were pooled from three separate experiments). (E) Sorted pDCs from LNs and spleens of naive BALB/c mice were pulsed with vehicle or rOpn for 24 h and then subjected to transmigrate in response to CCL19. Numbers of migrating PBS-treated pDCs (white bars) or rOpn- treated pDCs (black bars). Triplicate wells of pooled pDCs sorted from spleens and LNs (n = 8). Data are mean 6 SEM (n = 5–8 mice per group), one representative of five independent experi- ments (A–C) and from three independent experiments (E). *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.

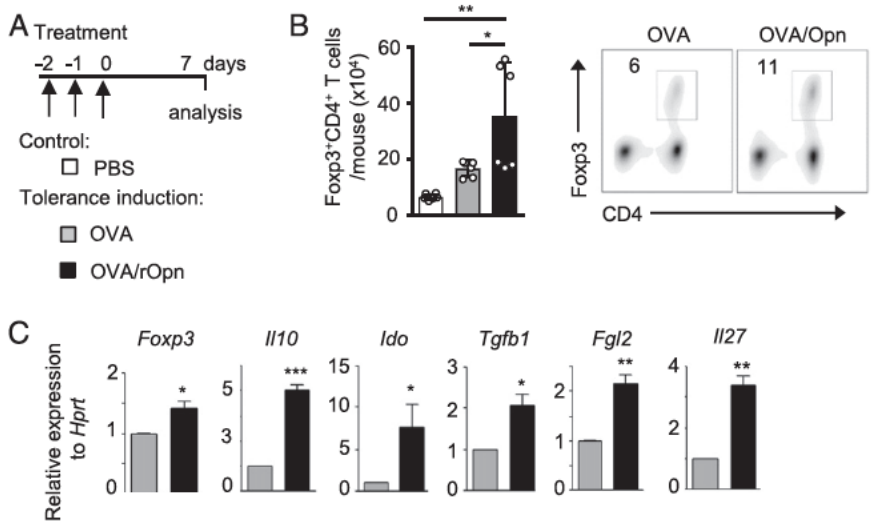


FIGURE 4. Opn administration with Ag promotes accumulation of Treg cells and immunoregulatory gene expression in dLNs. (A) Mice received three doses of endotoxin-free OVA for tolerance induc- tion, along with rOpn or PBS. Controls (without tolerance) were PBS-treated mice (white bars). (B) Percentages and numbers of CD3+CD4+Foxp3+ T cells quantified in dLNs by flow cytometry at day 7. (C) Relative expression to Hprt of immuno- regulatory genes in dLNs. cDNAs were pooled from three separate experiments. Data are mean 6 SEM (n = 6 mice per group), one representative of three inde- pendent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002.

OVA/alum	OVA aerosol
OVA/alum	OVA aerosol
OVA/alum	OVA aerosol
PBS/alum	OVA aerosol

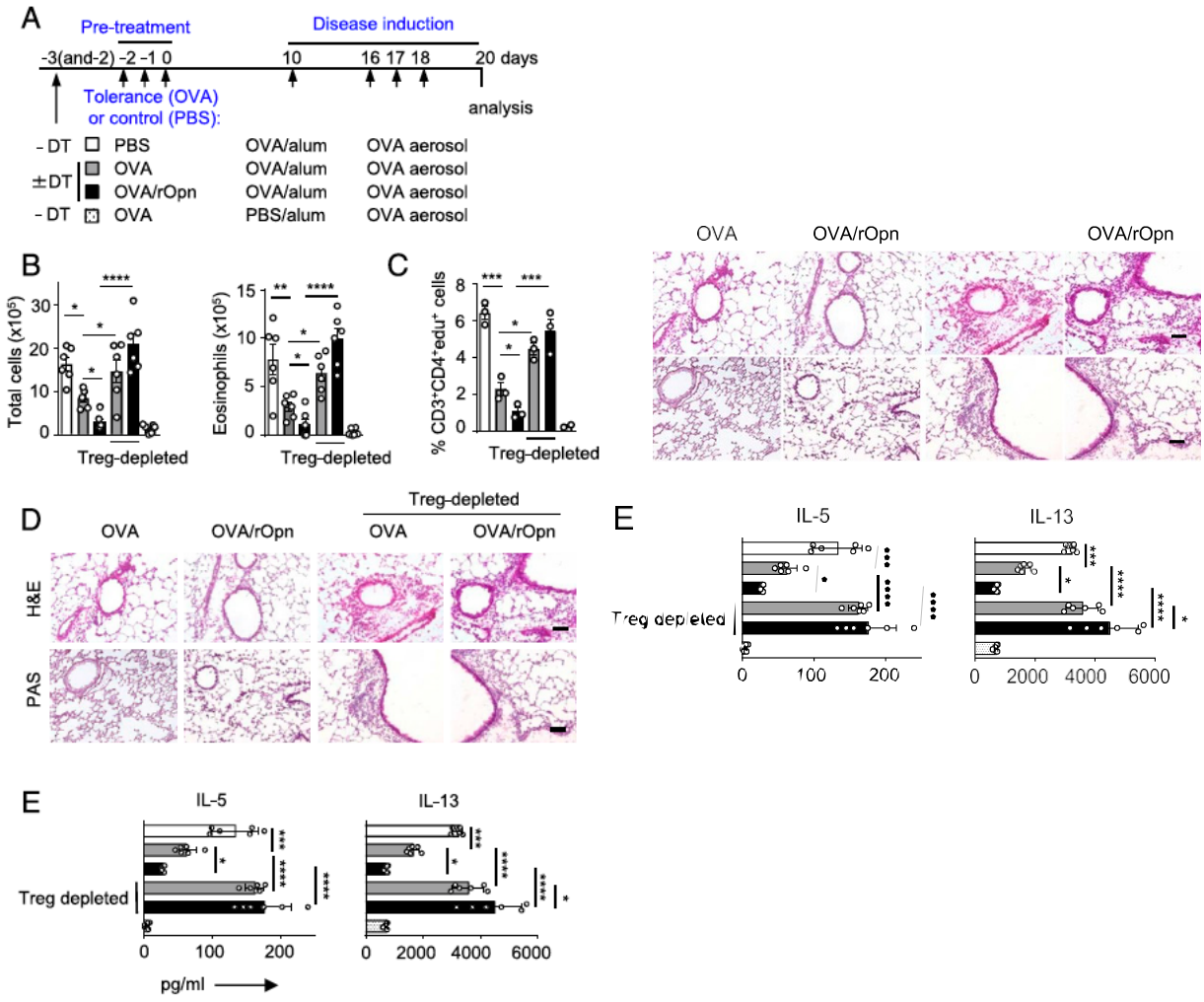


FIGURE 5. Opn-induced tolerance is Treg cell dependent. (A) Tolerance induction in DT-treated DEREG and control DEREG mice without DT injections. Controls (without tolerance) were DEREG mice pretreated withPBS, with subsequent allergy induction (white bars). (B) Eosinophil and total cell counts in BAL. (C) Numbers of $Edu^{+}CD3^{+}CD4^{+}$ T cells per 25 3 10^{4} OVA-stimulated dLN cells. (D) Lung inflammation and mucus se- cretion in H&E-stained (upper panels) and PAS-stained (lower panels) slides. Scale bars, 100 mm. (E) Levels of IL-5 and IL-13 in supernatants ofOVA-stimulated dLN cells. The same numbers of pooled mLN cells were cultured in three (C) or six (D) different wells per group. Data are mean 6 SEM (n = 6 mice per group), one representative of three independent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.

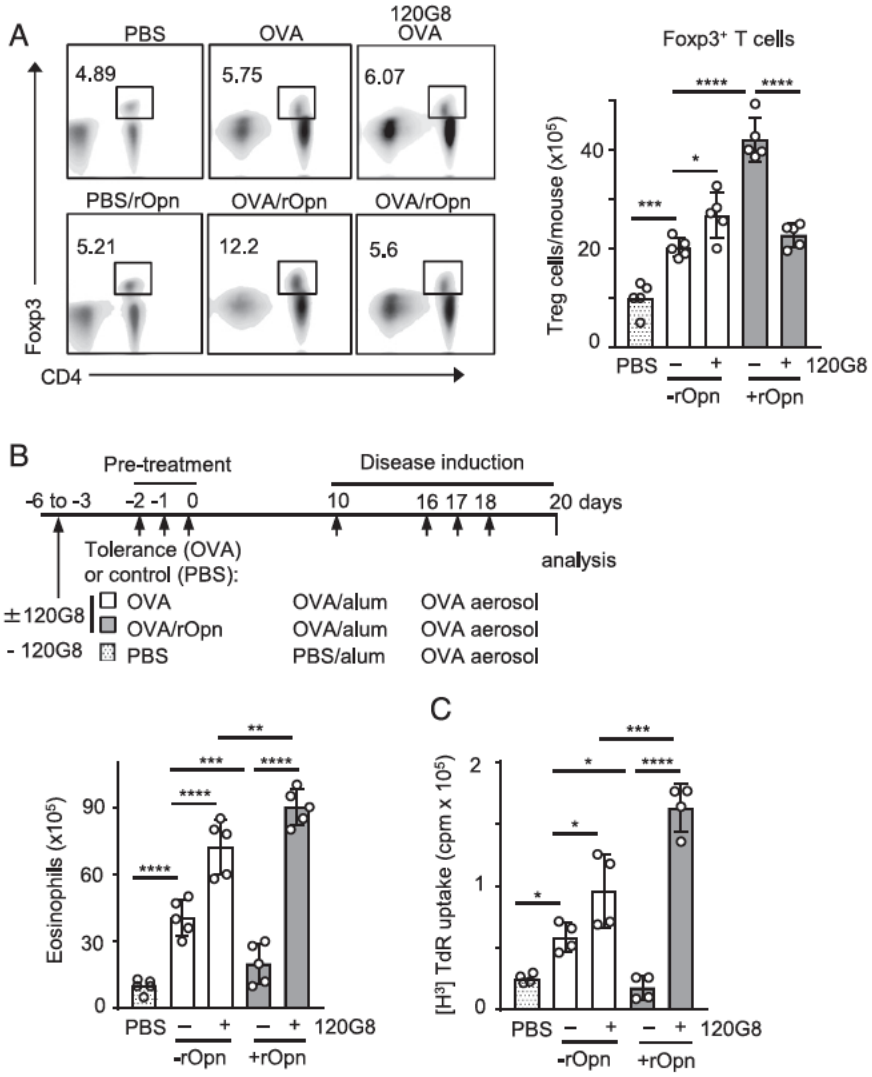


FIGURE 6. Opn-induced tolerance in- creases pDC numbers, favoring Treg cell accumulation. (A) Mice received four doses of 120G8 pDC-depleting Ab or Ig control i.p. on days 26, 25, 24, and 23, followed by tolerance induction for Treg cell generation, with or without rOpn ad- ministration. (B and C) Also, allergic air- way disease was subsequently induced on day 10. Control mice were pretreated with PBS (A) or sensitized with PBS (B and C). (A) Representative percentages in flowcytometric plots (left panel) and numbers of Treg cells (CD3+CD4+Foxp3+) accu- mulated in dLNs (right panel). (B) Eosin- ophil counts in BAL of allergic mice. (C) [3H]Thymidine incorporation in OVA- stimulated dLN cells. Quadruplicate wells of pooled mLN cells. Data are mean 6 SEM (n = 5 mice per group), one repre- sentative of three independent experi- ments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.

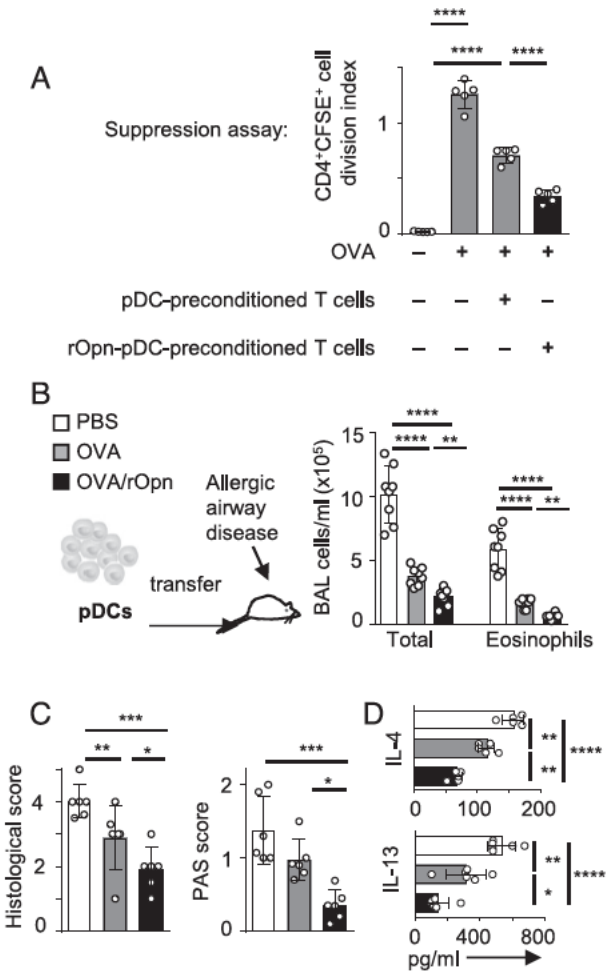


FIGURE 7. rOpn treatment of pDCs increases their suppressive activity against allergic airway disease. (A) [^{3}H]Thymidine incorporation in co-cultures of DO11.10 T cells preconditioned with $OVA_{323–339}$ and rOpn- treated pDCs with responder DO11.10 T cells. The same numbers of cells were cultured in five different wells per group. Data are mean 6 SEM, onerepresentative of three independent experiments. (B) Sorted pDCs pre- conditioned in vitro with PBS or OVA or OVA/rOpn were adoptively transferred to recipient mice before the induction of allergic airway in- flammation. Total and eosinophil cell count in BAL were evaluated (day 20). Histological assessment of lung inflammation (H&E scoring) and lungmucus production (PAS score) (C) and levels of IL-4 and IL-13 (D) in supernatants of OVA-stimulated mLN cells. Different wells of pooled mLN cells. Data are mean 6 SEM (n = 6–8 mice per group), one repre- sentative of three independent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.

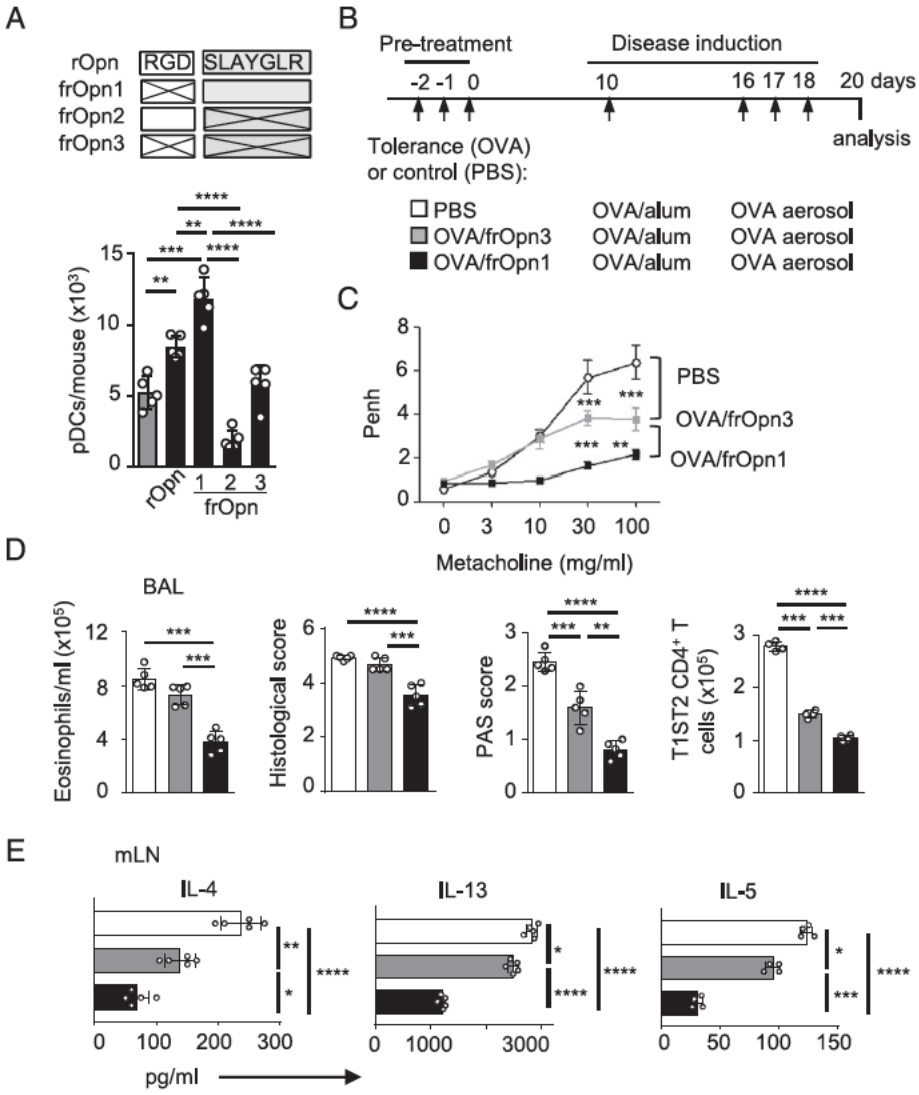


FIGURE 8. Opn SLAYGLR motifs boost tolerance by enhancing pDC recruitment to protect from al-lergy. (A) Numbers of dLN 7AAD^{2}CD11c^{+}B220^{+}PDCA1^{+}Siglec-H^{+} pDCs osinophil cell count in BAL, his- tological assessment of lung inflam- mation (H&E scoring) and lung mucus production (PAS score), and numbers of T1ST2+CD4+ TH2 cells in mLNs. after 40 h of tolerance induction with EndoGrade OVA i.n., along with rOpn, frOpn1, frOpn2, frOpn3, or PBS (as in Fig. 3A). (B) Mice were
treated with OVA/frOpn during tolerance induction before allergic air- way disease induction in (C)–(E). Control mice were pretreated with PBS, and allergy was subsequently induced (white bars). (C) AHR (depicted as Penh) at day 19. (D) Eosinophil cell count in BAL, histological assessment of lung inflammation (H&E scoring) and lung mucus production (PAS score), and numbers of T1ST2+CD4+ TH2 cells in mLNs. (E) Levels of IL-4, IL-5, and IL-13 in supernatants of OVA- stimulated mLN cells. The same numbers of pooled mLN cells were cultured in four to six different wells per group. Data are mean 6 SEM (n = 5, mice per group), one representative of three independent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.

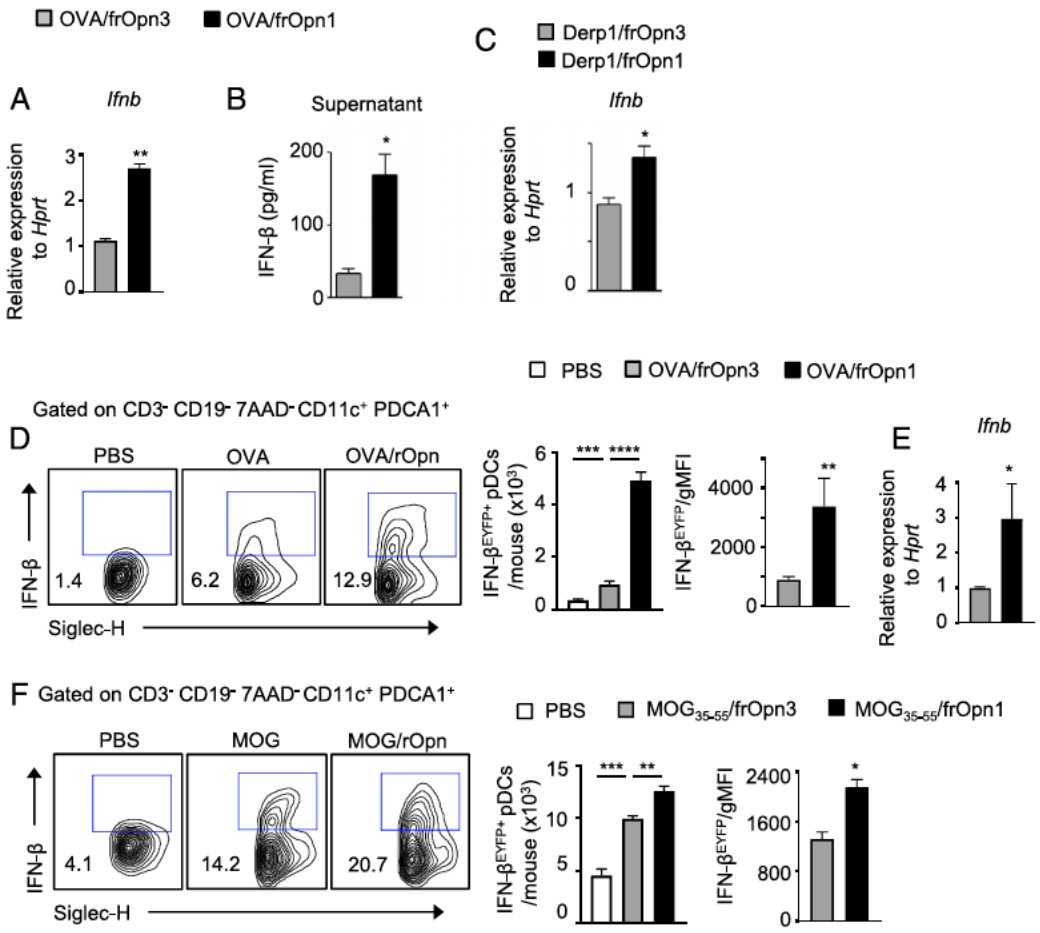


FIGURE 9. Opn SLAYGLR motif induces IFN-b production from pDCs. (A) Relative *Ifnb* expression to *Hprt* in pDCs conditioned in vitro with OVA/ frOpn1 or OVA/frOpn3. (B) Levels of IFN-b in the supernatants of the same cultures. (C) Relative *Ifnb* expression to *Hprt* in pDCs conditioned in vitro withDerp1/frOpn1 compared with Derp1/frOpn3. (D) Representative percentages in flow cytometric plots and numbers of $7AAD^{2}CD3^{2}CD19^{2}CD11c^{+}PDCA\text{-}1^{+}Siglec\text{-}H^{+}IFN\text{-}b^{EYFP+}$ pDCs in dLNs of PBS-treated or OVA-tolerized mice and geometrical MFI (gMFI) of IFN-b-expressing pDCs. (E) Relative *Ifnb*
expression to *Hprt* in pDCs isolated from dLNs of OVA/frOpn3- and OVA/frOpn1-tolerized mice. (F) Representative percentages in flow cytometric plots and numbers of total $7AAD^{2}CD3^{2}CD19^{2}CD11c^{+}PDCA\text{-}1^{+}Siglec\text{-}H^{+}IFN\text{-}b^{EYFP+}$ pDCs and gMFI of IFN-b–expressing pDCs in dLNs of mice treatedwith PBS or tolerized with $MOG_{35–55}$/frOpn3 or $MOG_{35–55}$/frOpn1. Mice were tolerized as in Fig. 3A. Data are mean 6 SEM (n = 6–8 mice per group),pooled data from three independent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.

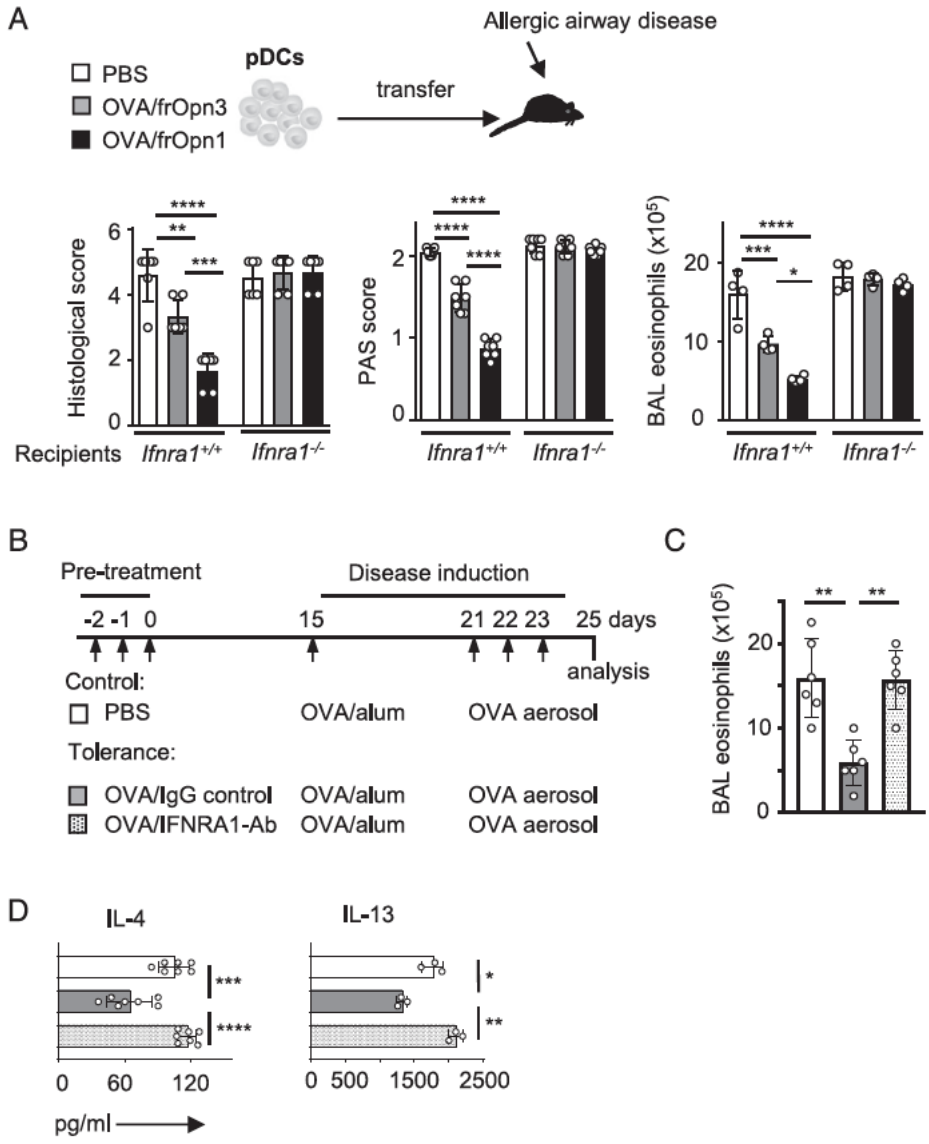
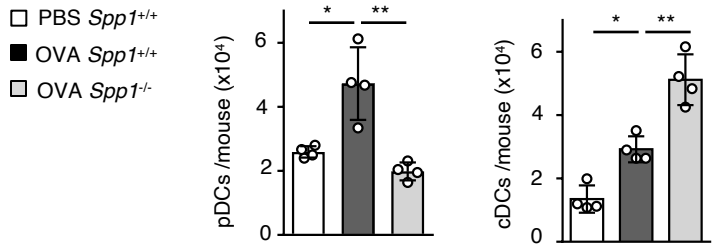
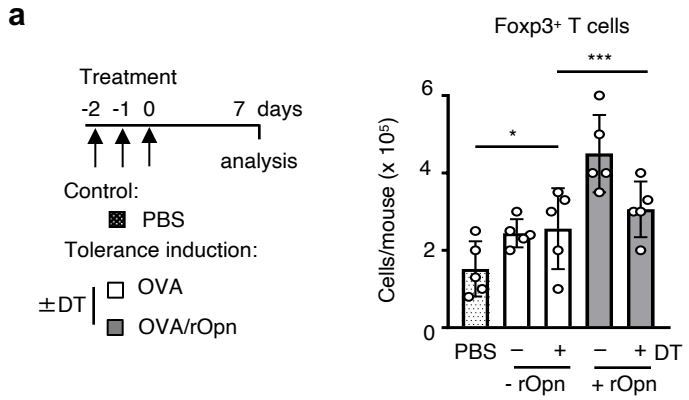


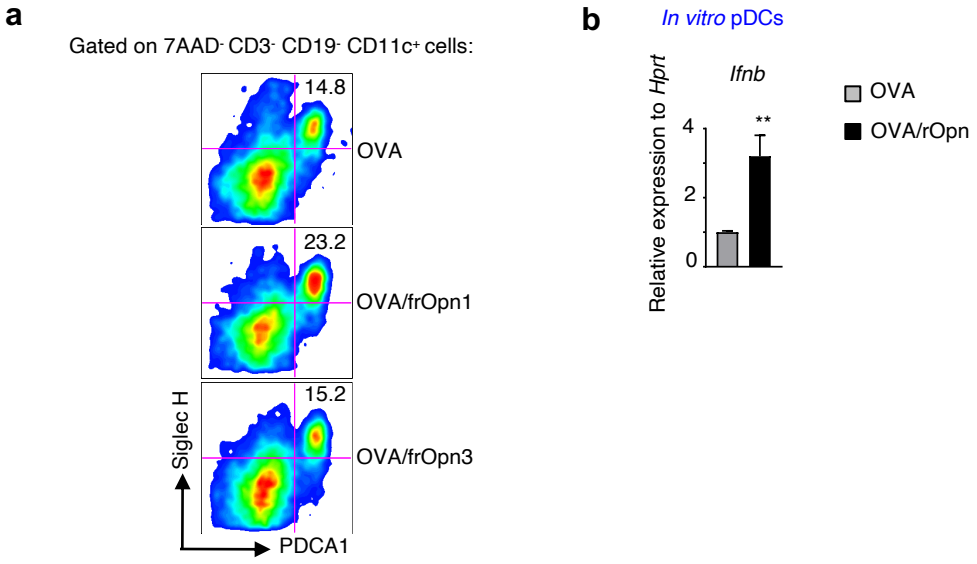
FIGURE 10. Opn/SLAYGLR-induced IFN-b in pDCs renders them protective against allergic disease. (A) Isolated *Ifnar1*$^{+/+}$ pDCs were primed in vitro withOVA and frOpn1 or frOpn3 or with PBS and adoptively transferred to *Ifnar1*$^{+/+}$ or *Ifnar*$^{2/2}$ mice. Histological scores for airway inflammation and goblet cellhyperplasia in lung sections stained with H&E and PAS, respectively, and number of eosinophils present in BAL. (B) OVA tolerance induction with the use of blocking Ab against IFNAR1 or isotype control prior to allergic airway disease induction. Control mice were pretreated with PBS prior to allergic airwaydisease induction (white bars). (C) Numbers of eosinophils in BAL and (D) levels of IL-4 and IL-13 in supernatants of OVA-stimulated dLN cells. The samenumbers of pooled mLN cells were cultured in three or seven different wells per group. Data are mean 6 SEM [n = 4 or 5 mice per group (A) and n = 6 mice per group (B–D)], one representative of three independent experiments. *p # 0.0332, **p # 0.0021, ***p # 0.0002, ****p , 0.0001.



Supplemental Figure 1. Opn increases the numbers of pDCs. Mice were pretreated with endotoxin-free OVA or PBS and analyzed after 40 hrs as in Figure 3a. Numbers of 7AAD^{-}CD3^{-}CD19^{-}CD11c^{+}CD11b^{-}B220^{+}PDCA-1^{+} pDCs in dLNs and 7AAD^{-}CD11c^{+}CD11b^{+}PDCA-1^{-} cDCs from dLNs of *Spp1*$^{+/+}$ or *Spp1*$^{-/-}$ mice. Values are expressed as mean ± SEM (n=4 mice per group), one representative experiment of three independent experiments.



Supplemental Figure 2. pDC deletion in BDCA2-DTR mice abrogates Treg cell accumulation in Opn-induced tolerance. For pDC depletion mice received diphtheria toxin (DT) on days -4, -3 followed by tolerance induction for Treg cell generation. Control mice were treated with PBS. Numbers of Treg cell ($CD3^+CD4^+Foxp3^+$) accumulation in dLNs with and without rOpn administration. Values are expressed as mean ± SEM (*n*=5 mice per group), one representative experiment of three independent experiments.



Supplemental Figure 3. Opn SLAYGR motif enhances pDC recruitment in dLNs and IFN-β production in pDCs. (A) Gating strategy and percentages of 7AAD^{-}CD3^{-}CD19^{-}CD11c^{+}PDCA-1^{+}Siglec-H^{+} pDCs in dLNs of tolerized mice treated with OVA or OVA/frOpn1 or OVA/frOpn3 (scrambled peptide: see Fig. 8A) for control. Mice were tolerized as in Fig. 3A. **(B)** Relative *Ifnb* expression to *Hprt* in pDCs conditioned *in vitro* with OVA and OVA/rOpn. Values are expressed as mean ± SEM (n=10), pooled data from three independent experiments.

Gene:	Sense (5'-3')	anti-sense (5'-3)
Ccl19	ATCTGAGCGATTCCAGTCA	TCTTCCGCATCATTAGCAC
Ccl21	TTCACGGTCCAACTCACA	CTCTTTCTTCTGACTCTCTA
Fgl2	ACAGCCGAGGACAGTAGA	CGGTTAGATTTGCCACTTTG
Ido	GGCTTCTTCCTCGTCTCTCT	ATTTCCTTGGCTTTCTCCAG
Tgfb1	CCTGGGTTGGAAGTGGA	TAGTAGACGATGGGCAGT
Ifnb	CCTATGGAGATGACGGAGAA	TGGAGAGCAGTTGAGGACAT

Supplemental Table 1: Primers used for real-time PCR.